













MICOPHENOLATE MOFETIL (2-(2-MECOXYBENZYL)BENZOPHENONE ACID)	MICOPHENOLATE MOFETIL	200 mg
UNITS/CONTAINER		

<b>Active Ingredients</b>	<b>Ingredient Name</b>	<b>Strength</b>
COPRO-CARBONATE (BISMUTH SUBGELATE)		
FIBR. 0.5% (L-LYSINE MONOCAPRYLATE)		
FIBR. 0.5% (L-LYSINE MONOCAPRYLATE)		
FIBR. 0.5% (L-LYSINE MONOCAPRYLATE)		
MICOPHENOLATE (2-(2-MECOXYBENZYL)BENZOPHENONE ACID)		
TITANIUM DIOXIDE (TITANIUM DIOXIDE)		

<b>Product Characteristics</b>	<b>Color</b>	<b>Strength</b>	<b>Unit</b>
	WHITE	200 mg	ea tablet
	Shape	CAPSULE	Size
	Color	White	Score
	Strength	200 mg	mg/tablet
	Container		

<b>Packaging</b>	<b>#</b>	<b>Drug Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
	1	NDC 074-074-074-2	100 X BOTTLE, PLASTIC		
	2	NDC 074-074-074-2	100 X BOTTLE, PLASTIC		

<b>Marketing Information</b>	<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
ANDA	ANDA019405		04/12/2010	

<b>MICOPHENOLATE MOFETIL</b>	<b>micophenolate mofetil capsule</b>
------------------------------	--------------------------------------

<b>Product Information</b>	<b>Product Type</b>	<b>HUMAN PRESCRIPTION DRUG</b>	<b>New Drug Order</b>	<b>NDA/ANDA/BLA</b>
	Route of Administration	ORAL	100A Schedule	

<b>Active Ingredients/Active Mixture</b>	<b>Ingredient Name</b>	<b>Strength</b>
MICOPHENOLATE (2-(2-MECOXYBENZYL)BENZOPHENONE ACID)		
UNITS/CONTAINER		
MICOPHENOLATE MOFETIL		
200 mg		

<b>Active Ingredients/Active Mixture</b>	<b>Ingredient Name</b>	<b>Strength</b>
COPRO-CARBONATE (BISMUTH SUBGELATE)		
FIBR. 0.5% (L-LYSINE MONOCAPRYLATE)		
FIBR. 0.5% (L-LYSINE MONOCAPRYLATE)		
FIBR. 0.5% (L-LYSINE MONOCAPRYLATE)		
MICOPHENOLATE (2-(2-MECOXYBENZYL)BENZOPHENONE ACID)		
TITANIUM DIOXIDE (TITANIUM DIOXIDE)		

<b>Product Characteristics</b>	<b>Color</b>	<b>Strength</b>	<b>Unit</b>
	WHITE	200 mg	ea tablet
	Shape	CAPSULE	Size
	Color	White	Score
	Strength	200 mg	mg/tablet
	Container		

<b>Packaging</b>	<b>#</b>	<b>Drug Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
	1	NDC 074-074-074-2	100 X BOTTLE, PLASTIC		
	2	NDC 074-074-074-2	100 X BOTTLE, PLASTIC		
	3	NDC 074-074-074-2	100 X BOTTLE, PLASTIC		

<b>Marketing Information</b>	<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
ANDA	ANDA019405		04/12/2010	

<b>Labeler</b>	Grenzauer LLC (20164773)
----------------	--------------------------

<b>Registrant</b>	Stedax Avicin Limited - NDA (108332a)
-------------------	---------------------------------------

<b>Establishment</b>	<b>Name</b>	<b>Address</b>	<b>RSP#</b>	<b>Business Operations</b>
----------------------	-------------	----------------	-------------	----------------------------

Stedax Avicin Limited				Business Operations
-----------------------	--	--	--	---------------------

Revised: 9/2010

Grenzauer LLC